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- Prophylactic and therapeutic agent for bone diseases comprising di- or tripeptide derivative as active ingredient.
- (I): Use of a compound represented by the following general formula

wherein R1 represents a straight or branched acyl group having 2 to 10 carbon atoms, a branched cyclic or bridged cyclic alkyloxycarbonyl group having 4 to 15 carbon atoms, a substituted or unsubstituted benzyloxycarbonyl group, a 2,2,2-trichloroethyloxycarbonyl group, a 2-(trimethylsilyl)-ethyloxycarbonyl group, a p-toluenesulfonyl group, an o-nitrophenylsulfenyl group, a diphenylphosphonothioyl group, a triphenylmethyl group, a 2benzoyl-1-methylvinyl group or a 4-phenylbutanoyl group;

R2 represents a hydrogen atom or forms a phthaloyl group together with R1;

R³ represents an isobutyl group, an n-butyl group or an isopropyl group;

R4 represents an n-butyl group or an isobutyl group;

X represents a direct bond, a methionine residue, a leucine residue or a norleucine residue; and H at the C-terminal means that the carboxyl group is reduced in the form of the aldehyde;

as an active ingredient for the preparation of a pharmaceutical composition useful in preventing or treating malignant hypercalcemia, bone Paget's disease or osteoporosis.

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Y	JOURNAL OF ENZYME I vol. 3, no. 3, 1990 pages 195 - 201 SASAKI ET AL 'Inhib tripeptidyl aldehyd cathepsins' * See the whole doc	, HARWOOD, ENGLAND itory effect of di- and es on calpains and	1-4	A61K37/02 C07K5/08
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	The present search report has be			
Ti	Place of search HE HAGUE	Date of completion of the nearch 19 FEBRUARY 1993		KORSNER S.E.
X : partic Y : partic docum A : techn	ATEGORY OF CITED DOCUMEN cularly relevant if taken alone cularly relevant if combined with another ment of the same category sological background written disclosure nather secument	E : earlier patent doc after the filing da	ument, but publi te the application other reasons	shed on, or

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EUROPEAN SEARCH REPORT

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